## Abstract of the Disclosure

A compound of the formula II

5 wherein

one of R<sup>1</sup> and R<sup>2</sup> is halo and the other is H or halo;

 $R^3$  is  $C_1\text{-}C_4$  straight or branched chain, optionally fluorinated, alkyl;

R⁴ is H; or

R<sup>3</sup> together with R<sup>4</sup> and the adjoining backbone carbon defines:

a spiro- $C_5$ - $C_7$  cycloalkyl, optionally substituted with 1 to 3 substituents selected from halo, hydroxyl,  $C_1$ - $C_4$  alkyl or  $C_1$ - $C_4$  haloalkyl; or optionally bridged with a methylene group; or

a C<sub>4</sub>-C<sub>6</sub> saturated heterocycle having a hetero atom selected from

O, NRa, S, S(=O)<sub>2</sub>; where Ra is H, C<sub>1</sub>-C<sub>4</sub> alkyl or CH<sub>3</sub>C(=O);

15 R<sup>5</sup> is independently selected from H or methyl;

E is -C(=O)-, -S(=O)<sub>m</sub>-, -NR $^5$ S(=O)<sub>m</sub>-, -NR $^5$ C(=O)-, -OC(=O)-,

R<sup>6</sup> is a stable, optionally substituted, monocyclic or bicyclic, carbocycle or hetorocycle; m is independently 0,1 or 2;

are inhibitors of cathepsin K and useful in the treatment or prophylaxis of osteoporosis.